mg. per lb. Alphatocopherol (Vit. E) 200 mg. per lb. Ascorbic Acid (Vit. C) 200 mg. per lb."

RESULTS OF INVESTIGATION: The article was shipped unlabeled as described above and after its receipt by the dealer at Slayton, Minn., the tins containing the article were labeled with the above-mentioned labels.

LIBELED: 3-28-60, Dist. Minn.

CHARGE: 502(a)—the label of the article, while held for sale, contained false and misleading representations that the article was an adequate and effective treatment and preventive of all diseases in poultry, hogs, calves, lambs, dogs, cats, and mink; and 502(l)—the article was a drug composed in part of penicillin and streptomycin sulfate and it was not from a batch with respect to which a certificate or release had been issued pursuant to 507 in that certification of the article under the above-mentioned label had not been obtained.

DISPOSITION: 5-11-60. Consent—claimed by Tracy L. Hafner, t/a Tracy Hafner Slayton Drug Store, and released for relabeling.

DRUGS IN VIOLATION OF PRESCRIPTION LABELING REQUIREMENTS

6204. Seconal Sodium capsules and amphetamine tablets. (F.D.C. No. 43585. S. No. 66-005 P.)

QUANTITY: 50,000 tablets of amphetamine and 1,000 capsules of Seconal Sodium at Stamford, Conn.

SHIPPED: On or about 10-8-59, from New Jersey to Connecticut, by Charles W. Christiansen, also known as Charlie Benjamin.

LIBELED: On or about 10-8-59, Dist. Conn.

CHARGE: 502(b)—the articles failed to bear labels containing (1) the name and place of business of the manufacturer, packer, or distributor and (2) an accurate statement of the quantity of contents in terms of weight, measure, or numerical count; 502(d)—the *Seconal Sodium capsules* contained a chemical derivative of barbituric acid, and their label failed to bear the name, and quantity, or proportion of such derivative and in juxtaposition therewith the statement "Warning—May be habit forming"; 502(e)(1)—the labels of the articles failed to bear the common or usual names of the articles; 502(f)(1)—the labeling of the articles failed to bear adequate directions for use and they were not exempt from such requirement since the articles were in possession of persons who were not regularly and lawfully engaged in the manufacture, transportation, storage or wholesale distribution of prescription drugs and since the articles were not to be dispensed as required by 503(b)(1); and 503(b)(4)—the articles were subject to 503(b)(1) and their labeling failed to bear the mandatory statement "Caution: Federal law prohibits dispensing without prescription."

DISPOSITION: 6-28-60. Default—destruction.

DRUGS AND DEVICES ACTIONABLE BECAUSE OF FAILURE TO BEAR ADEQUATE DIRECTIONS OR WARNING STATEMENTS*

6205. Various drugs. (F.D.C. No. 44590. S. Nos. 19-841 R, 19-843 R, 19-846 R, 19-851/4 R, 19-856 R, 19-858/9 R, 19-861 R, 19-864 R, 19-868/70 R, 19-872 R, 19-875 R, 19-877 R, 19-881 R, 21-067/71 R, 21-073/4 R, 21-077 R, 21-079 R, 21-083 R, 21-087/8 R, 21-093/4 R, 21-097 R, 21-099/100 R, 21-103 R, 21-107 R, 21-110/11 R, 21-113 R, 21-117/19 R, 21-122 R.)

*See also Nos. 6202, 6204.